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In the claims:

1. (Currently Amended) A method of administering to a subject in need thereof an effective amount of a cisplatin active agent, said method comprising:

administering to said host subject said effective amount of a cisplatin active agent in conjunction with an amount of a cisplatin toxicity reducing agent compound selected from TK-211; TK-295; TK-516; TK-523; TK-363; TK-204; TK-5145 and TK-5175 effective to reduce toxicity of said cisplatin active agent.

- 2. (Currently Amended) The method according to Claim 1, wherein said cisplatin active agent and cisplatin toxicity reducing agent said compound are administered at the same time.
- 3. (**Currently Amended**)The method according to Claim 2, wherein said cisplatin active agent and cisplatin toxicity reducing agent said compound are administered as separate formulations.
- 4. (**Currently Amended**)The method according to Claim 2, wherein said cisplatin active agent and cisplatin toxicity reducing agent said compound are administered in a single formulation.
- 5. (**Currently Amended**)The method according to Claim 1, wherein said cisplatin active agent and cisplatin toxicity reducing agent said compound are administered sequentially.
- 6. (**Currently Amended**)The method according to Claim 5, wherein said cisplatin active agent is administered prior to said eisplatin toxicity reducing agent compound.

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7. (**Currently Amended**) The method according to Claim 5, wherein said cisplatin active agent is administered after said cisplatin toxicity reducing agent compound.

8. (**Currently Amended**) The method according to Claim 1, wherein the amount of said **cisplatin toxicity reducing agent compound** is not more than about the amount of said cisplatin **active agent**.

9.-16. (**CANCELLED**)

17. (Currently Amended) A method of treating a host suffering from a cellular proliferative disease condition ovarian cancer in a patient, said method comprising:

administering to said host patient said an effective amount of a cisplatin active agent cisplatin effective to treat said cancer in conjunction with an amount of a cisplatin toxicity reducing agent TK-211 effective to reduce toxicity of said cisplatin active agent so that said host is treated for said cellular proliferative disease condition.

- 18. (Currently Amended) The method according to Claim 17, wherein said cisplatin active agent and cisplatin toxicity reducing agent said TK-211 are administered at the same time.
- 19. (**Currently Amended**) The method according to Claim 18, wherein said cisplatin active agent and cisplatin toxicity reducing agent said TK-211 are administered as separate formulations.
- 20. (Currently Amended) The method according to Claim 18, wherein said cisplatin active agent and cisplatin toxicity reducing agent said TK-211 are administered in a single formulation.

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21. (**Currently Amended**) The method according to Claim 17, wherein said cisplatin active agent and cisplatin toxicity reducing agent said TK-211 are administered sequentially.

- 22. (**Currently Amended**) The method according to Claim 21, wherein said cisplatin-active agent is administered prior to said cisplatin toxicity reducing agent TK-211.
- 23. (**Currently Amended**) The method according to Claim 21, wherein said cisplatin active agent is administered after said cisplatin toxicity reducing agent <u>TK-211</u>.
- 24. (**Currently Amended**) The method according to Claim 17, wherein the amount of said **cisplatin toxicity reducing agent** <u>TK-211</u> is not more than about the amount of said cisplatin <u>active agent</u>.

Claims 25 -30. (CANCELLED)